Applicant: Aladar Szalay et al. Attorney's Docket No.: 3800002.00055/4804US

Serial No.: 10/516,785 Preliminary Amendment with RCE

Filed : June 27, 2005

AMENDMENTS TO THE CLAIMS:

Please amend claims 1, 13, 15 and 23 as follows. This listing of claims replaces all prior versions, and listings of claims, in the application.

LISTING OF CLAIMS:

(Currently Amended) A method for detecting wounded or inflamed tissue 1. inside of a subject, comprising:

systemically administering to a subject in whom the presence or absence of a wounded tissue or inflamed tissue or a disease associated therewith is to be detected, a bacterium, wherein:

the bacterium is detectable in the subject;

the bacterium replicates in the subject;

the bacterium is not pathogenic to the subject and is recognized by the immune system of the subject;

the bacterium is not targeted; and

monitoring the subject to detect the accumulation of the bacterium at or in a wounded tissue or inflamed tissue within inside of the subject, whereby, wherein detection of the accumulation indicates the location of wounded tissue or inflamed tissue inside of within the subject, thereby detecting wounded or inflamed tissue inside of the subject.

- 2. (Previously Presented) The method of claim 1, wherein the bacterium encodes a protein(s) for the therapy of the detected wounded or inflamed tissue.
- 3. (Previously Presented) The method of claim 1, wherein the bacterium encodes a luminescent or fluorescent protein.
- 4. (Previously Presented) The method of claim 1, wherein the bacterium encodes a luciferase, red fluorescent protein or green fluorescent protein.
- 5. (Previously Presented) The method of claim 4, wherein the bacterium encodes a luciferase and a protein(s) for the production of a substrate for a luciferase.
- 6. (Previously Presented) The method of claim 1, wherein the bacterium encodes a protein that induces a signal detectable by magnetic resonance imaging (MRI) or that binds to contrasting agent, chromophore or a ligand.
 - 7. (Cancelled).
 - 8. (Cancelled).

2 of 14

Applicant: Aladar Szalay et al. Attorney's Docket No.: 3800002.00055/4804US

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9. (Previously Presented) The method of claim 1, wherein the bacterium is selected among an attenuated Salmonella typhimurium, an attenuated Vibrio cholerae, an attenuated *Listeria monocytogenes* and *E. coli*.

- 10. (Cancelled).
- 11. (Cancelled).
- 12. (Previously Presented) The method of claim 2, wherein the protein for the therapy is an enzyme that causes cell death or an enzyme that causes the digestion of debris.
- 13. (Currently Amended) The method of claim 2, wherein the subject for in whom the presence or absence of wounded tissue or inflamed tissue is detected has a disease selected among endocarditis, pericarditis, inflammatory bowel disease, low back pain (herniated nucleus pulposis), temporal arteritis, polyarteritis nodosa and an arthritic disease.
- 14. (Previously Presented) The method of claim 2, wherein the subject for whom the presence or absence of wounded tissue or inflamed tissue is detected has an atherosclerotic disease.
- 15. (Withdrawn, Currently Amended) The method of claim 2, wherein the subject for whom the presence or absence of wounded tissue or inflamed tissue is detected has a disease that is selected among coronary artery disease, peripheral vascular disease and cerebral artery disease.
- (Previously presented) The method of claim 1, wherein the monitoring is 16. performed by magnetic resonance imaging (MRI).
 - 17. (Cancelled).
- 18. (Previously Presented) The method of claim 2, wherein the bacterium contains an inducible promoter that regulates the expression of the therapeutic protein.
 - 19. (Cancelled).
 - 20. (Cancelled).
 - 21. (Previously presented) The method of claim 2, wherein:

the disease is an atherosclerotic disease; and

the therapeutic protein is selected from among a lipase, protease, lysozyme, proapoptotic factor and PPAR-agonist.

(Previously Presented) The method of claim 1, further comprising: 22. administering a therapeutic agent for the therapy of a wounded tissue, inflamed tissue or a disease associated therewith.

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23. (Currently Amended) The method of claim 1, wherein the bacterium is administered intravenously, intraperitoneally, subcutaneously, or intramuscularly, topically or intradermally.